



Continuous, non-invasive measurement of the haemodynamic response to submaximal exercise in patients with diabetes mellitus. Evidence of impaired cardiac reserve and peripheral vascular response

Deepa Joshi, Ajay Shiwalkar, Maurice Cross, et al.

Heart published online October 22, 2009
doi: 10.1136/hrt.2009.177113

Updated information and services can be found at:
<http://heart.bmj.com/content/early/2009/10/22/hrt.2009.177113>

These include:

Open Access

This manuscript is Open Access.

P<P

Published online October 22, 2009 in advance of the print journal.

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

Advance online articles have been peer reviewed and accepted for publication but have not yet appeared in the paper journal (edited, typeset versions may be posted when available prior to final publication). Advance online articles are citable and establish publication priority; they are indexed by PubMed from initial publication. Citations to Advance online articles must include the digital object identifier (DOIs) and date of initial publication.

To order reprints of this article go to:
<http://heart.bmj.com/cgi/reprintform>

To subscribe to *Heart* go to:
<http://heart.bmj.com/subscriptions>

Continuous, non-invasive measurement of the haemodynamic response to submaximal exercise in patients with diabetes mellitus. Evidence of impaired cardiac reserve and peripheral vascular response

Deepa Joshi^{*}, Ajay Shiwalkar^{*}, Maurice R Cross[§], Sanjay K Sharma[#], Ashish Vachhani^{*}, Chaitanya Dutt^{*§}.

Word Count: 3141

Department where the work is carried out: Discovery Biology, Torrent Research Centre, Village Bhat, Gandhinagar 382428, Gujarat, India and Clinical Pharmacology Unit, Veeda Clinical Research, Shivalik Plaza – A, Near IIM, Ambawadi Ahmedabad 380015, Gujarat, India

Authors Affiliations: ^{*}Torrent Research Centre, Village Bhat, Gandhinagar 382428, Gujarat, India. [§]Veeda Clinical Research, Old Convent of Notre Dame, 119 Looseleigh Lane, Derriford, Plymouth PL65HH, UK. [#]Veeda Clinical Research, Shivalik Plaza – A, Near IIM, Ambawadi Ahmedabad 380015, Gujarat, India

[§]Corresponding author:

Dr. Chaitanya Dutt

Torrent Research Center

P.O. Bhat, Dist.Gandhinagar -382428

Gujarat, India.

Tel: +91-79-23969100 Fax: +91-79-23969135

Email: cdutt@torrentpharma.com

EXCLUSIVE LICENCE

"The Corresponding Author has the right to grant on behalf of all authors and does grant on behalf of all authors, an exclusive license (or non exclusive for government employees) on a worldwide basis to the BMJ Publishing Group Ltd and its Licensees to permit this article to be published in HEART editions and any other BMJ PGL products to exploit all subsidiary rights as set out in our licence <http://heart.bmjournals.com/ifora/licence.pdf> "

ABSTRACT

Background: Reduced exercise capacity in diabetics has been attributed to limitations in cardiac function and microvascular dysfunction leading to impaired supply and nutritive perfusion to exercising muscles. **Objective:** To study changes in cardiac function, and microvascular utilization during exercise, in diabetic individuals compared to age-matched controls. **Methods:** Diabetics with HbA1c<8 (n=31), diabetics with HbA1c≥8 (n=38), and age-matched non diabetic controls (n=32) performed exercise at 50W for 10 minutes followed by recovery, with continuous monitoring of cardiac function by impedance cardiography and regional flow and oxygen saturation by laser Doppler and white light spectroscopy. **Results:** In the diabetics, cardiac reserve during exercise and cardiac overshoot during recovery are significantly reduced due to reduction in capacity to increase stroke volume. Regional flow to the exercising muscle is reduced and there is disproportionately greater desaturation of the regional flow as well. Abnormalities in cardiac function and regional perfusion are related to the severity of diabetes. **Conclusion:** This study shows that cardiac response to exercise is attenuated significantly in diabetic individuals. Simultaneously, there is impairment in the regional distribution. These changes could be the harbinger of reduced exercise capacity in diabetics.

Keywords: diabetes, exercise, impedance cardiography, muscle, oxygen

BACKGROUND

Clinical studies have shown that limitation in exercise capacity is a strong predictor of cardiovascular and all cause mortality in diabetic patients with heart failure. Underlying pathophysiology is multifactorial and involves alterations in ventricular-vascular coupling consisting of cardiac changes on one hand and compromised distribution at the regional microvasculature, on the other. If such changes can be detected early (Stage A) in those who are at risk of progressing to heart failure, it may help slow down progression by more aggressive management of the problem.

Though there is fair body of evidence on the mechanism underlying the decline in exercise tolerance and despite the recognition of its prognostic significance, monitoring microvascular changes underlying altered skeletal muscle exercise capacity and mapping it dynamically against cardiac response capacity has not been attempted. Monitoring of regional alterations in the microcirculation in patients is restricted by the invasive character of methodology and limitations to its adaptation to record continuous and dynamic change in the exercising muscle. Assessing cardiac function during exercise poses a special challenge as conventional echocardiographic measures including strain analysis, can only be done at rest or at the end of exercise. Therefore dynamic response to exercise which is a function of continuous adjustment of pre-load (Frank-Starling) and after-load as well as direct cardiac adrenergic influence, designed to optimize the inotropic and chronotropic responses, cannot be captured by these methods.

Impedance cardiography (ICG) has been validated to gather ambulatory physiological recordings and was adapted to monitor cardiac function online during exercise.[1, 2] For monitoring regional microcirculation this was combined with another method wherein laser Doppler shift caused by the movement of erythrocytes in the detected laser light is used to determine arteriolar flow. With the same probe, white light is introduced into the tissue and oxygen saturation of hemoglobin was determined from light absorbed by the hemoglobin in mixed capillary venous blood at a predetermined depth from the skin.[3] This study was therefore aimed at investigating the sequential changes during mild exercise and recovery in cardiac function and regional microcirculation of exercising muscle in diabetes by a method which allows continuous dynamic monitoring, in order to capture early cardiac and vascular correlates, which could ultimately progress to impaired exercise tolerance. Submaximal exercise is more relevant as it is equivalent to 6-minute walk test, used to evaluate exercise capacity in patients with marked LV dysfunction,[4] in contrast, VO_2 max may be compromised due to reduced patient motivation, poor peripheral blood flow and impaired skeletal muscle metabolism with early development of acidosis besides, limitation in cardiac output, correlates only modestly with this test.[5]

STUDY PROTOCOL

Subjects

The investigation conforms to the principles outlined in the Declaration of Helsinki.[6] The study was approved by the Institutional Review Board of Torrent Research Centre and Veeda Clinical Research (Ahmedabad, India). After signing the written informed consent, 127 subjects were screened. The inclusion criteria were subjects with confirmed diabetes as ascertained by history, clinical examination and laboratory investigations, who had no history of breathlessness on exertion or limitation in exercise capacity, and age-matched controls who were non-diabetic. Subjects with arrhythmia, significant peripheral vascular disease, retinopathy, insulin dependence and those with blood pressure (BP)>180/100 mmHg, were excluded from the study.

101 Subjects were included in the study and all completed the protocol. Data from diabetics with Haemoglobin A1c (HbA1c) <8 (n=31), diabetics with HbA1c ≥ 8 (n=38), and age-matched normal controls (n=32) were included in the analysis.

Experimental procedure

At Veeda Clinical Research, after their breakfast subjects had a brief physical examination, blood samples were collected for estimation of HbA1c, vascular cell adhesion molecule (VCAM), N-terminal brain natriuretic propeptide (NTProBNP), and urine samples for creatinine and microalbumin estimation. Oxygen to See (O2C, LEA Medizintechnik, Germany) probe LF3 incorporating Doppler (830nm) and near infra-red light spectrum (500-850nm), which allows measurement at skeletal muscle post-capillary venular site, was placed on the belly of the gastrocnemius on one leg. ICG (Bioz, Cardiodynamics, USA) and electrocardiogram (ECG-Mac5000, GE Healthcare, Germany) probes were placed on the chest and blood pressure (BP) cuff on the arm.

After calibration, continuous measurements were obtained at baseline in resting supine position for 10 minutes followed by upright position for 10 minutes and then acclimatization on the bicycle ergometer (eBike, GE Healthcare, Germany) for 10 minutes. Exercise was for duration of 10 minutes, at constant load of 50 Watts, on the electrically braked ergometer, varying rpm to maintain the constant load, which is equivalent to 4-7 METS. This level of exercise was chosen to simulate a 6-minute walk test commonly used to evaluate patients with heart failure in order to determine if it was adequate to evoke peripheral vascular response to differentiate the diabetic from non-diabetic subjects. At the end of exercise period, subjects were asked to lie supine again for a recovery period of at least 15 minutes. ECG was recorded throughout and BP every two minutes. The same protocol was repeated in 10 subjects to assess the repeatability of the measurements. The entire study was completed over a period of four months.

Data acquisition and processing

Data from O2C & ICG were acquired for five different phases viz. resting supine baseline, standing, sitting on ergometer, during exercise & recovery after exercise. ICG measurements (collected every 1 minute) & O2C measurements (collected every 2 seconds was also averaged for 1 minute) were classified & synchronized for different phases of the protocol. Data were cleaned for machine error and artifacts (random and during transition from one phase of protocol to next) for all the measured variables. For O2C parameters, flow response to exercise was measured as area under curve (AUC) and Oxygen saturation (SO₂) as area over the curve (AOC) of percent change at each time point during exercise, from mean of sitting. In addition, slope to achieve maximum velocity in the microcirculation, was computed. ICG parameters considered for analysis were cardiac output (CO), stroke volume (SV) and systolic time interval. Systolic time interval was obtained by summing the Pre-ejection period (PEP) and Left ventricular ejection time (LVET) obtained from ICG. Response during recovery was measured as AUC of the data points recorded during recovery phase. Subjects who completed the exercise phase of 10 minutes were included for analysis on exercise data (Table-1), and for analysis on recovery data, only those who completed exercise followed by 15 minutes of recovery were included (diabetics with HbA1c <8 n=28, diabetics with HbA1c ≥ 8 n=35 and age-matched normal controls n=32). To calculate Oxygen debt for each subject, the value of mean SO₂ at resting supine baseline, multiplied by the same period of time i.e. 15 minutes, was subtracted from AUC over 15 minutes of recovery.[7] AUC has been calculated using WinNonlin (Version-5.2).

Statistical analysis

Data are expressed as mean \pm SD and graphically represented as mean \pm SEM. Statistical analysis has been performed using SAS (Version-9.1). Demographic data has been tested to see differences among 3 groups using ANOVA followed by Post-Hoc Dunnett t-tests (two-sided). Repeatability was determined by two-sided paired t-test on AUC values (derived from % change during exercise from sitting baseline) of the O₂C parameters, SO₂ and flow and ICG parameter, CO. Proportions of concomitant medications were subjected to Fisher's exact test. CO, SV, time for downward inflection, and AUC or AOC of percent change from baseline, during exercise and for recovery period, was compared between groups by one-sided unpaired t-test. In absence of normality, data were subjected to non-parametric Wilcoxon test. Percent change in systolic time interval over time, of Normal, Diabetics with HbA_{1c}<8 and Diabetics with HbA_{1c}≥8, were compared by repeated measures analysis of variance (RMANOVA). Statistical significance was defined as 'p' value \leq 0.05.

RESULTS

Demography

Demographic profile of the subjects included in the study is shown in Table 1. Medications, sulphonylurea secretagogues, thiazolidinedione sensitizers, antihypertensives ACE inhibitors and ARBs and loop diuretics, besides statin and beta blockers (Table 1), taken by the subjects did not influence the measurements made during the study. Statins were analysed separately and did not show an effect on the cardiac response to exercise when comparing the two diabetic groups. Similarly, the diabetic group with $HbA1c \geq 8$, showed a greater attenuation of the cardiac response to exercise as compared to the diabetic group with $HbA1c < 8$. This was probably not secondary to attenuation of the response due to beta blockers as fewer diabetics with $HbA1c \geq 8$ were taking beta blockers as compared to those with $HbA1c < 8$.

Cardiac response during exercise

Resting CO was lowest in normal individuals and although statistically not significant, it was slightly higher in the diabetic groups with $HbA1c < 8$ and ≥ 8 . In response to exercise, there was a rise in both SV and CO in all groups. However, the capacity to maximally increase CO in response to exercise, a measure of the cardiac reserve, was significantly ($p < 0.001$) reduced in the diabetic patients as compared to normal subjects (Fig 1A). This could be ascribed to a smaller rise in SV during exercise in diabetic patients, which was significantly reduced ($p < 0.002$) in the more severely diabetic group (Fig 1B). Following initial rise, the time to downward inflection towards reduction of SV, an indicator of inadequate filling secondary to increasing heart rate, occurred significantly ($p < 0.05$ N vs < 8 ; $p < 0.001$ N vs ≥ 8) earlier, and at lower heart rates ($p < 0.05$ N vs diabetic groups) in the diabetic groups as compared to normal (Fig 2A and 2B). Lengthening of the systolic time interval was observed in the diabetic group with $HbA1c \geq 8$, in that there was greater compensatory lengthening of systolic time interval depicted here as less fall in the systolic time for a given stroke output during exercise as compared to normal ($p < 0.05$) (Fig 2C).

Cardiac response during recovery

During the recovery phase after exercise, the CO remained elevated and did not return to resting baseline. The increase however was attenuated in diabetics, more so in the diabetics with $HbA1c \geq 8$ ($p < 0.05$) (Fig 3A). There was no difference in the heart rate response during recovery between the different groups.

Skeletal muscle perfusion during exercise

Regional changes in perfusion in the muscle as revealed by local measurement of flow are shown in figure 4A. There was a significant ($p<0.05$) reduction in AUC of % change from baseline in diabetic individuals with $HbA1c \geq 8$ (1024.7 ± 780.3) as compared to normal subjects (1433.4 ± 1055.9). Diabetics with $HbA1c < 8$, on the other hand, did not show a significant difference, however, to compensate, flow was slightly increased (1541.1 ± 1648.1) as compared to normal (Fig 4A). This was accompanied by a reduction in oxygen saturation in both the diabetic groups as compared to normal (Fig 4B), AOC in normal (-77.2 ± 170.8) was significantly ($p<0.05$) more compared to diabetics with $HbA1c < 8$ (-125.3 ± 236.1) and diabetics with $HbA1c \geq 8$ (-165.6 ± 247.6). This was offset, in the diabetics with $HbA1c \geq 8$, by increased velocity of blood in the microcirculation, which showed a significant increase ($p<0.05$) in the slope to achieve maximal velocity (slope= 3.05 ± 2.59), as compared to the normal subjects (slope= 2.08 ± 1.65).

Skeletal muscle perfusion during recovery

Regional flow to the exercising muscle normally remains elevated, over resting levels, even after stopping exercise, during recovery. Compared to normal subjects, in diabetics, augmentation of flow was more (Fig 3B), in order to compensate for the reduced CO response during recovery.

Oxygen debt as determined from the diminished SO_2 during recovery, over the resting baseline, was significantly increased in the diabetic subjects as compared to normal (Fig 3C) ($p<0.05$). Despite the increase in flow, desaturation of the local flow over the resting values persisted, however, it was least in the normal subjects, as evident by highest AUC of oxygen saturation during recovery. In the diabetic group, SO_2 reduced significantly in both mild and severely diabetic patients (Fig 3D).

Repeatability

Data from ten subjects who repeated the exercise protocol showed that AUC of % change during exercise from sitting position to exercise did not show any significant difference in SO_2 (507 ± 158 initial vs 625 ± 126 repeat, $p>0.05$), flow (3400 ± 867 initial vs 3166 ± 996 repeat, $p>0.05$) and in CO (75.6 ± 13.3 initial vs 75.7 ± 13.8 repeat, $p>0.05$). Hence, the precision of the relative measurements before and during exercise were reasonably accurate for acceptance of O₂C and ICG.

DISCUSSION

This study characterizes the hemodynamic response to exercise in diabetic as compared to normal individuals, by dynamic recording of cardiac function as well as simultaneous monitoring of regional perfusion of the muscle. These findings demonstrate that cardiac response even to submaximal exercise is attenuated significantly in diabetic individuals. Simultaneously, there is an attempt to maintain regional microvascular perfusion by a compensatory augmentation of regional flow to the exercising muscle. Despite this compensation, there is greater desaturation of the supply reaching the working muscle in diabetics compared with normal individuals. Furthermore, the reduction in cardiac reserve as well as regional oxygen delivery during exercise, were found to be related to the severity of diabetes.

Reduced exercise capacity in diabetics has been reported in non insulin dependent diabetes mellitus (NIDDM) subjects without coronary artery disease [8] and a correlation between left ventricular dysfunction and impaired exercise capacity has been demonstrated in diabetics in several studies.[9] Our study confirms that the SV reserve is the key parameter that is compromised during exercise in asymptomatic diabetic individuals. This is evident even with the mild exercise equivalent to a 6-minute walk test when measured dynamically in ambulatory subjects who are not in failure. The association of poor glycemic control with worse cardiac and peripheral dysfunction in this study is consistent with a previous large study in asymptomatic type-2 diabetes.[10] Other studies have shown that A1c has an inverse correlation with maximum oxygen uptake,[11] work capacity,[12] and exercise duration.[13]

Perturbation of cardiac function independent of coronary artery disease and BP in diabetes may result from an interplay of a wide variety of progressively dysfunctional autoregulatory mechanisms. Diastolic abnormalities like delayed onset of ventricular relaxation,[14] and systolic function abnormalities of the ventricular muscular pump characterized by abnormal systolic time intervals and by increased brain natriuretic peptide (BNP) levels, as observed in this study, precede reduced haemodynamic pump performance with severe systolic and diastolic abnormalities. Limitations on cardiac reserve and cardiac contractile function thus restrict the supply from meeting the demand imposed by exercise. In addition, microcirculatory deficits coupled with altered energetics limit the ability to meet the demand in exercising skeletal muscle in diabetics.[15,16] Endothelial dysfunction and functional alterations of the microcirculation, on the other hand, precede morphological changes and contribute to an increased vascular permeability [17] and an impaired endothelial vasodilatory response. There are several reports that insulin also increases sympathetic nervous system activity which can determine the nutritive flow distribution.[18] Diabetes with associated insulin resistance and autonomic dysfunction could lead to further exacerbation of the maldistribution locally to fulfill the nutritive flow required to meet the metabolic demand during exercise. These changes acting in conjunction, induce dysfunction with resultant reductions in flow and O₂ delivery at rest and during exercise.[19] A perturbation of intramuscular homeostasis in response to exercise challenge was also observed in this study, which has the potential to contribute to premature muscular fatigue and progress to reduced exercise tolerance in diabetic individuals.

During the recovery period, a shift from carbohydrate to fat as a fuel substrate occurs to prevent further depletion of muscle glycogen stores, leading to oxygen consumption remaining elevated above resting levels after exercise, called oxygen debt, because the oxygen cost of fat catabolism is greater than that of carbohydrate catabolism.[20,21] In order to meet this requirement, marked increase in stroke volume during early recovery with overshoot appears to be facilitated by both an immediate afterload reduction and a relatively slow decrease in cardiac sympathetic

stimulation. Accumulation of a large oxygen debt during exercise has been reported in heart failure patients.[22] In this study, overshoot of cardiac output was observed during recovery, though lower in diabetics and so was the mixed venous SO_2 , suggesting more severe muscle under perfusion and greater O_2 -deficit and regional ischemia in diabetic individuals. Possible mechanism could be a persistence of sympathetic activity which serves to increase vascular resistance in non-working tissues during exercise, which the local metabolic factors and endothelium-derived substances such as nitric oxide and prostaglandins are unable to override, as they are deficient in diabetic state. Increased renin–angiotensin–endothelin system activity may also contribute, at least in part.

Mediators of the changes underlying impaired exercise response in patients with type-2 diabetes are multiple. Limited exercise tolerance has been associated with poor glycemic control.[8, 10-12, 23] Secondary to this, glycosylation leading to formation of AGE, initiate a proinflammatory and procoagulant state and may impair the function of a number of proteins leading to cardiac, vascular or endothelial dysfunction.[24, 25] We have found that diabetes-induced myocardial dysfunction in diabetic hypertensive rat model was reversed by reducing the AGE burden consequent to treatment with AGE-breaker.[26]

Previous reports of ICG in diabetics have been limited and restricted to measurements at rest. One study in diabetic individuals revealed that thoracic fluid content increased despite improvement in contractile indices after treatment with Rosiglitazone.[27] In another study, no difference was detected in cardiac index at rest, between healthy individuals and newly diagnosed diabetics, [28] indicating that possibly exercise provocation would be necessary to bring out the subtle differences in the hemodynamics. Insights on the control of muscle blood flow and oxygen consumption during exercise have, in past, relied on measurements of bulk flow in large conduit arteries to the exercising limbs precluding analysis of local perfusion and oxygenation mismatch. However comparing changes under resting, dynamic exercise and recovery after exercise with a combination of NIRS, laser Doppler and impedance cardiography to expose sub-clinical differences in diabetic individuals has not been attempted previously to the best of our knowledge. Repeatability of the measurements showed reasonable consistency between measurements made on both O₂C and ICG parameters.

Potential limitations of our study are, the mixed diabetic population studied, which could have incipient complications like neuropathy, peripheral vascular disease and coronary artery disease, which could be influencing the study parameters. For diabetic subjects without any signs of heart failure, reduction in exercise capacity secondary to reduction in cardiac output may be evident only during maximal exercise and could be of prognostic significance. We also do not have tissue-doppler data to correlate limitation in increasing stroke volume to diastolic dysfunction.

CONCLUSION

In summary, this cross-sectional study in diabetic population showed that by combining the assessment of dynamic cardiac response during exercise with peripheral flow distribution and nutritive perfusion, sub-clinical differences in hemodynamic response were detected in diabetic individuals, hitherto not reported to the best of our knowledge. Longitudinal follow-up of these patients and a comparison of dynamic cardiac response and that of peripheral flow and nutritive distribution, as described, with the traditional measures of exercise capacity which have been studied longitudinally over time and shown to have prognostic significance would provide an objective and sensitive measure of cardiac and microvascular correlates of the composite

hemodynamic function and indicate the suitability of such diagnostic measures for early detection of a population at risk of progressing to Heart Failure.[5]

ACKNOWLEDGMENTS: We are grateful to Dr. Jinesh Shah, Mr. Anand Chettiyar and Mr. Nilesh Badgujar for assistance in conduct and data management of the study and Ms. Shikha Kumar for editorial assistance.

COMPETING INTEREST: The authors DJ, AS, AV and CD are employees of Torrent Pharmaceutical Limited, Ahmedabad and MRS and SKK are employees of Veeda UK and India respectively.

FUNDING: The study was funded by Torrent Pharmaceutical Limited.

Figure Legends

Figure 1 (A) Cardiac reserve during exercise and (B) Stroke volume reserve during exercise, the key parameter limiting increase in cardiac output.

Figure 2 (A) Time to downward inflection of stroke volume response during exercise demonstrating that, with increasingly poor glycemic control there is increasing limitation in the capacity to augment stroke volume (B) Heart Rate at Downward Inflection in Stroke Volume During Exercise. Increase in CO occurs as a function of increase in heart rate with decreasing oxygen pulse delivered to muscles (C) Reduction in systolic time interval compared to baseline indicating longer systole and compromised diastole in diabetics.

Figure 3 (A) Difference in AUC of cardiac output during recovery from resting baseline: Cardiac Output overshoot during recovery (B) Increase in regional flow over baseline during recovery: Difference in AUC during recovery from resting baseline (C) Oxygen debt: Difference in AUC during recovery from resting baseline (D) Oxygen Saturation (AUC) in muscle flow during recovery.

Figure 4 (A) Regional muscle flow during exercise and (B) oxygen saturation in blood supplied to exercising muscle. Note that in diabetics with reasonable control increase in flow is not able to contain desaturation. Diabetics with poor control show limited capacity to augment flow and run up a steeper oxygen debt with a late attempt at correction which could be due to local factors overriding systemic vascular tone.

REFERENCES

1. Thomas SH, Crowther A. Impedance cardiography during exercise in patients with coronary heart disease. *Eur Heart J* 1993;**14**(2):150-9.
2. Greenberg BH, Hermann DD, Pranulis MF, *et al.* Reproducibility of impedance cardiography hemodynamic measures in clinically stable heart failure patients. *Congest Heart Fail* 2000;**6**(2):74-80.
3. Beckert S, Witte MB, Koenigsrainer A, *et al.* The Impact of the Micro-Lightguide O2C for the Quantification of Tissue Ischemia in Diabetic Foot Ulcers. *Diabetes Care* 2004;**27**:2863-7.
4. Bittner V, Weiner DH, Yusuf S, *et al.* Prediction of mortality and morbidity with a 6-minute walk test in patients with left ventricular dysfunction: SOLVD Investigators. *JAMA* 1993;**270**:1702-7.
5. Davies LC, Wensel R, Georgiadou P, *et al.* Enhanced prognostic value from cardiopulmonary exercise testing in chronic heart failure by non-linear analysis: oxygen uptake efficiency slope. *Eur Heart J* 2006;**27**:684-90.
6. Rickham PP. Human Experimentation code of ethics of the world medical association. Declaration of Helsinki. *BMJ* 1964;**ii**:177.
7. Aliverti A, Dellacà RL, Lotti P, *et al.* Influence of expiratory flow-limitation during exercise on systemic oxygen delivery in humans. *Eur J Appl Physiol* 2005;**95**(2-3):229-42.
8. Estacio RO, Regensteiner JG, Wolfel EE, *et al.* The association between diabetic complications and exercise capacity in NIDDM patients. *Diabetes Care* 1998;**21**(2):291-5.
9. Fang ZY, Sharman J, Prins JB, *et al.* Determinants of exercise capacity in patients with Type 2 Diabetes. *Diabetes Care* 2005;**28**(7):1643-8.
10. Demir I, Ermis C, Altunbas H, *et al.* Serum HbA1c levels and exercise capacity in diabetic patients. *Jpn Heart J* 2001;**42**(5):607-16.
11. Vanninen E, Uusitupa M, Siitonen O, *et al.* Habitual physical activity, aerobic capacity and metabolic control in patients with newly-diagnosed type 2 (non-insulin-dependent) diabetes mellitus: effect of 1-year diet and exercise intervention. *Diabetologia* 1992;**35**(4):340-6.
12. Barkai L, Peja M, Vamosi I. Physical work capacity in diabetic children and adolescents with and without cardiovascular autonomic dysfunction. *Diabet Med* 1996;**13**(3):254-8.
13. Baum VC, Levitsky LL, Englander RM. Abnormal cardiac function after exercise in insulin-dependent diabetic children and adolescents. *Diabetes Care* 1987;**10**(3):319-23.
14. De Keulenaer GW, Brutsaert DL. Systolic and diastolic heart failure: different phenotypes of the same disease? *Eur J Heart Fail* 2007;**9**(2):136-43.
15. Scheuermann-Freestone M, Madsen PL, Manners D, *et al.* Abnormal cardiac and skeletal muscle energy metabolism in patients with Type 2 Diabetes. *Circulation* 2003;**107**(24):3040-6.

16. Wu YW, Hsu CL, Wang SS, *et al.* Impaired exercise capacity in diabetic patients after coronary bypass surgery: effects of diastolic and endothelial function. *Cardiology* 2008;**110**:191-8.
17. Kingwell BA, Bradley SJ, Formosa BA, *et al.* Type 2 diabetic individuals have impaired leg blood flow responses to exercise: role of endothelium-dependent vasodilation. *Diabetes Care* 2003;**26**(3):899-904.
18. Vollenweider P, Randin D, Tappy L, *et al.* Impaired insulin-induced sympathetic neural activation and vasodilation in skeletal muscle in obese humans. *J Clin Invest* 1994;**93**(6):2365-71.
19. Creager MA, Luescher TF, Cosentino F, *et al.* Diabetes and Vascular Disease. Pathophysiology, Clinical Consequences, and Medical Therapy: Part I. *Circulation* 2003;**108**(12):1527-32.
20. Wong T, Harber V. Lower excess postexercise oxygen consumption and altered growth hormone and cortisol responses to exercise in obese men. *J Clin Endocrinol Metab* 2006;**91**(2):678-86.
21. Kano H, Koike A, Yajima T, *et al.* Mechanism of overshoot in cardiac function during recovery from submaximal exercise in man. *Chest* 1999; **116**(4):868-73.
22. Mitchell SH, Steele NP, Leclerc KM, *et al.* Oxygen cost of exercise is increased in heart failure after accounting for recovery costs. *Chest* 2003;**124**(2):572-79.
23. Fang ZY, Prins JB, Marwick TH. Diabetic cardiomyopathy: evidence, mechanisms, and therapeutic implications. *Endocr Rev* 2004;**25**(4):543-67.
24. Wautier JL, Schmidt AM. Protein glycation: A firm link to endothelial cell dysfunction. *Circ Res* 2004;**95**:233-38.
25. Kelley DE, He J, Menshikova EV, *et al.* Dysfunction of Mitochondria in Human Skeletal Muscle in Type 2 Diabetes. *Diabetes* 2002;**51**(10):2944-50.
26. Pathak P, Gupta R, Chaudhari A, *et al.* TRC4149 a novel Advanced Glycation End Product breaker improves hemodynamics status in diabetic spontaneously hypertensive rats. *Eur J Med Res* 2008;**13**:388-98.
27. Ferdowsali K, Tadlaoui K, Nguyen E, *et al.* Rosiglitazone induced changes in the hemodynamic profile of patients assessed by impedance cardiography. *Circulation* 2006;**114**:II_881 Abstract.
28. Loimaala A, Groundstroem K, Majahalme S, *et al.* Impaired myocardial function in newly onset type 2 diabetes associates with arterial stiffness. *Eur J Echocardiogr* 2006;**7**(5):341-47.

Table 1. Demographic profile of subjects (Mean \pm SD or [#]Median \pm interquartile range)

	Normal (n=32)	HbA1C < 8 (n=31)	HbA1C \geq8 (n=38)
Age (y)	48.8 (5.3)	50.6 (7.3)	50.3 (4.6)
BMI (kg/m ²) [§]	23.81* \pm 3.4	26.42 \pm 2.8	26.57 \pm 3.8
Systolic Blood Pressure (mmHg)	116 \pm 10	120 \pm 12	117 \pm 11
Diastolic Blood Pressure (mmHg)	80 \pm 9	80 \pm 7	78 \pm 9
Cardiac Output (L/ml)	4.96 \pm 0.95	5.10 \pm 0.91	5.19 \pm 0.76
Heart Rate	77 \pm 11	80 \pm 14	80 \pm 10
NT - proBNP (pg/ml) [#]	31 (14.5 - 51.5)	35.1 (11.5 - 62.7)	27.6 (14.6 - 55.7)
VCAM (ng/ml) [#]	326.8 (186 - 647)	343.8 (171 - 617)	416.9 (253 - 611)
Diabetic Duration (y) [#]	-	4 (3 - 10)	5 (3 - 10)
Microalbuminuria (mg/L)	18.99 \pm 44.93	17.85 \pm 36.35	41.05 \pm 88.11
eGFR (ml/min/1.73 m ²)	85 \pm 12	80 \pm 16	90 \pm 17
Concomitant medication			
Secretagogues	-	70	73
Sensitizers	-	73	76
Beta blockers [§]	-	33	5**
Antihypertensives	-	50	34
Diuretics	-	7	3
Statins	-	8.3	0

*p<0.01 N vs <8, N vs \geq 8; **p<0.05 <8 vs \geq 8[§]Does not alter and contribute to Diabetic vs Non-diabetic comparison for various ICG & O2C variables





A



B



C



D



